

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
4 April 2002 (04.04.2002)

PCT

(10) International Publication Number
WO 02/26921 A1

(51) International Patent Classification⁷: C11D 1/02, 1/62,
3/24, 3/36, 7/36, D06L 1/02, 1/04, 1/22

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(21) International Application Number: PCT/US01/29812

(22) International Filing Date:
25 September 2001 (25.09.2001)

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(25) Filing Language: English

(81) Designated States (national): AU, CA, CN, ES, GB, JP,
KR, MX, SG.

(26) Publication Language: English

(84) Designated States (regional): European patent (AT, BE,
CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC,
NL, PT, SE, TR).

(30) Priority Data:
60/235,516 26 September 2000 (26.09.2000) US

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Published:

— with international search report

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

(54) Title: PHOSPHATE FLUOROSURFACTANTS FOR USE IN CARBON DIOXIDE

(57) Abstract: A method of removing water from a composition of matter comprises contacting a first composition of matter comprising water with a second composition of matter comprising: (1) at least one surfactant comprising at least one phosphate group and (2) a solvent comprising carbon dioxide, wherein at least a portion of the surfactant is soluble in the solvent, such that the at least one surfactant removes at least a portion of the water from the first composition

PHOSPHATE FLUOROSURFACTANTS FOR USE IN CARBON DIOXIDE

Cross-Reference to Related Applications

The present application claims priority to Provisional Application Serial No. 60/235,516 filed September 26, 2000, the disclosure of which is incorporated herein by reference in its entirety.

Field of the Invention

The present invention generally relates to surfactants that exhibit solubility in carbon dioxide, and systems utilizing the same.

Background of the Invention

The use of carbon dioxide as a clean, abundant, and tunable solvent is potentially environmentally beneficial, and accordingly it is being investigated in a number of applications including, for example, cleaning protocols, coatings, and polymer production and processing. See e.g., Wells, S. L.; DeSimone, J. *Angew. Chem. Int. Ed.* **2001**, *40*, 518. Notwithstanding the above potential benefits, carbon dioxide is often limited in that many materials such as water exhibit limited solubility therein.

In response to these possible solubility limitations, fluorosurfactants have been developed to potentially assist in the dispersion of water in carbon dioxide. See e.g., Harrison, K.; Goveas, J.; Johnston, K. P.; O'Rear, E. A. *Langmuir* **1994**, *10*, 3536; Johnston, K. P.; Harrison, K. L.; Clarke, M. J.; Howdle, S. M.;

Heitz, M. P.; Bright, F. V.; Carlier, C.; Randolph, T. W. *Science* **1996**, *271*, 624, Eastoe, J.; Bayazit, Z.; Martel, S.; Steytler, D. C.; Hennen, R. K. *Langmuir* **1996**, *12*, 1423, Eastoe, J.; Cazelles, B. M. H.; Steytler, D. C.; Holmes, J. H.; Pitt, A. R.; Wear, T. J.; Heenan, R. K. *Langmuir* **1997**, *13*, 6980, Zielinski, R. G.; Kline, S. R.; Kaler, E. W.; Rosov, N. *Langmuir* **1997**, *13*, 3934, Eastoe, J.; Downer, A.; Paul, A.; Steytler, D. C.; Rumsey, E.; Penfold, J.; Heenan, R. K. *Phys. Chem. Chem. Phys.* **2000**, *2*, 5235, Lee Jr., C. T.; Bhargava, P.; Johnston, K. P. *J. Phys. Chem. B* **2000**, *104*, 4448, Lee Jr., C. T.; Johnston, K. P.; Dai, H. J.; Cochran, H. D.; Melnichenko, Y. B.; Wignall, G. D. *J. Phys. Chem. B* **2001**, *105*, 3540, and Liu, Z.-T.; Erkey, C. *Langmuir* **2001**, *17*, 274. Accordingly, water-in-carbon dioxide (W/C) microemulsions containing appreciable water quantities have been achieved, allowing for their use in a number of applications such as nanoparticle synthesis, organic reactions, voltammetric measurements, and enzymatic conversions. See e.g., Holmes, J. D.; Bhargava, P. A.; Korgel, B. A.; Johnston, K. P. *Langmuir* **1999**, *15*, 6613, Ji, M.; Chen, X.; Wai, C. M.; Fulton, J. L. *J. Am. Chem. Soc.* **1999**, *121*, 2631, Jacobson, G. B.; Lee Jr., C. T.; Johnston, K. P. *J. Org. Chem.* **1999**, *64*, 1201, Ode, H.; Hunt, F.; Kithara, S.; Way, C. M. *Anal. Chem.* **2000**, *72*, 4738, Lee, D.; Hutchison, J. C.; Demimonde, J. M.; Murray, R. M. *J. Am. Chem. Soc.* **2001**, *123*, 8406, Holmes, J. D.; Settler, D. C.; Rees, G. D.; Robinson, B. H. *Langmuir* **1998**, *14*, 6371, and Kane, M. A.; Baker, G. A.; Pander, S.; Bright, F. V. *Langmuir* **2000**, *16*, 4901. W/C systems have also been the subject of computational treatments. See e.g., Satanicelli, S.; Cui, S. T.; Cummings, P. T.; Cochran, H. D. *Langmuir* **1999**, *15*, 5188.

Nonetheless, in spite of any advantages of these systems, there remains a need in the art for surfactants and systems employing the same that allow for improved volatilization of various materials in carbon dioxide, such as, for example, water.

Summary of the Invention

In one aspect, the invention provides a method of removing water from a first composition of matter. The method comprises contacting a first composition of matter comprising water with a second composition of matter comprising: (1) at least one surfactant comprising at least one phosphate group and (2) a solvent comprising carbon dioxide, wherein at least a portion of the surfactant is soluble in the solvent, such that the at least one surfactant removes at least a portion of the water from the first composition.

In another aspect, the invention provides a method of applying a surfactant to a substrate. The method comprises providing a composition of matter comprising at least one surfactant comprising (1) at least one phosphate group and (2) a solvent comprising carbon dioxide, wherein at least a portion of the surfactant is soluble in the solvent; and applying the composition of matter onto a substrate such that the carbon dioxide separates from the surfactant and wherein the surfactant coats the substrate.

In another aspect, the invention provides compositions of matter comprising (1) at least one surfactant comprising at least one phosphate group and (2) a solvent comprising carbon dioxide.

These and other aspects of the invention are described in greater detail herein.

Brief Description of the Drawings

FIG. 1 illustrates a ^{31}P NMR spectra of [2-(*F*-hexyl)ethyl] dimorpholinophosphate before (top) and after (bottom) being solubilized in carbon dioxide.

FIG. 2 illustrates cloud point curves for various phosphate-containing surfactants.

FIG. 3 illustrates cloud point curves for various phosphate-containing surfactants.

FIG. 4 illustrates cloud point curves for various phosphate-containing anionic surfactants.

FIG. 5 illustrates cloud point curves for various phosphate-containing anionic surfactants.

FIG. 6 illustrates water uptake for various phosphate-containing anionic surfactants.

FIG. 7 illustrates water uptake for various phosphate-containing anionic surfactants.

FIG. 8 illustrates water uptake for various phosphate-containing anionic surfactants.

FIG. 9 illustrates water uptake for various phosphate-containing anionic surfactants.

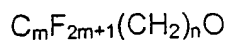
Detailed Description of the Preferred Embodiments

The invention will be described below with respect to its preferred embodiments, drawings, and examples. It should be appreciated that these do not serve to limit the scope of the invention, but instead illustrate the scope of the invention.

In one aspect, the invention relates to a composition of matter. The composition of matter comprises: (1) at least one surfactant comprising at least one phosphate group and (2) a solvent comprising carbon dioxide.

The surfactant may be illustrated in a number of embodiments. In various embodiments, the surfactant comprises at least one fluorocarbon group. For the purposes of the invention, the at least one fluorocarbon group is considered to have an affinity for carbon dioxide, i.e., the at least one fluorocarbon group is "CO₂-philic". Preferably, the at least one fluorocarbon group comprises a hydrogen spacer attached to an oxygen atom which is, in turn, attached to the phosphate group.

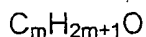
In various preferred embodiments, the at least one fluorocarbon group is of the formula:



wherein m ranges from 1 to 24 and n ranges from 1 to 24.

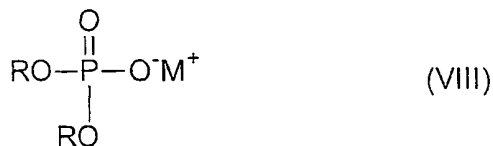
In other embodiments, the surfactant may further comprise a hydrocarbon-containing group attached to the phosphate group. In accordance with the invention, the hydrocarbon-containing group is, in certain embodiments, "CO₂-phobic", does not have affinity for carbon dioxide. Nonetheless, it should be appreciated that when present in branched form, the hydrocarbon-containing group may be useful for dispersing charged surfactants in CO₂. See Eastoe *et al. Journal of the American Chemical Society*; 2001; 123(5); 988-989.

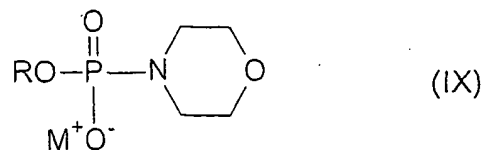
In a preferred embodiment, the hydrocarbon-containing chain is of the formula:



wherein m ranges from 1 to 24.

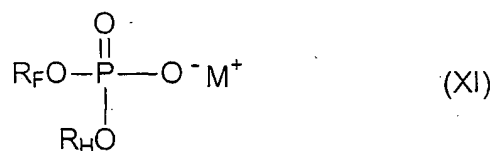
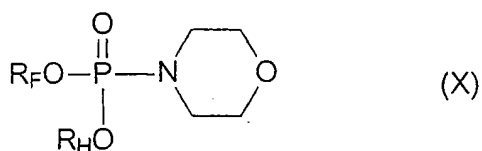
In various embodiment, the surfactant may be present as an anionic surfactant. Non-limiting examples of anionic surfactants are of the formulas (VIII) and (IX) as well as others set forth in detail hereinbelow:

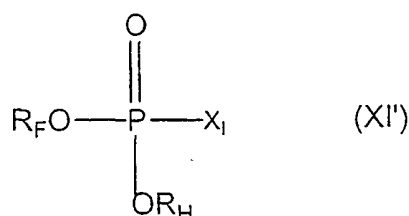




wherein R is a branched or straight chained hydrocarbon (e.g., $\text{C}_n\text{H}_{2n+1}$ wherein n ranges from 1 to 24) or hydrocarbon/fluorocarbon group and M is a counteranion such as, for example, K^+ , Na^+ , or NH_4^+ .

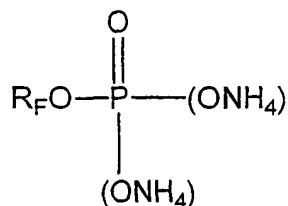
In certain embodiments, the surfactants may be present in the form of hybrid surfactants. For the purposes of the invention, the term "hybrid surfactant" is defined as a surfactant having at least one fluorocarbon group and at least one hydrocarbon-containing group. Examples of these are of the formulas (X), (XI), and (XI'): As set forth below, the hybrid surfactants also encompass anionic surfactants in various embodiments.





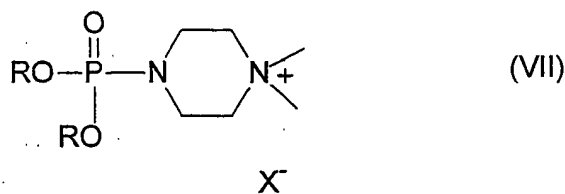
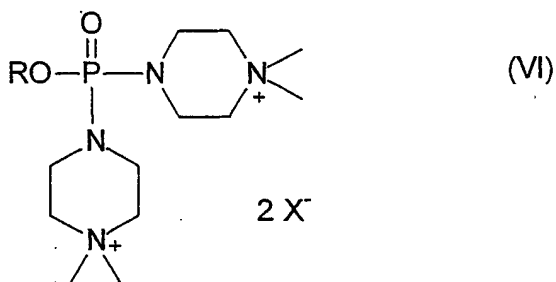
wherein R_H may be independently selected and is a branched or straight-chained hydrocarbon group (e.g., $\text{C}_n\text{H}_{2n+1}$ wherein n ranges from 1 to 24). R_H is preferably C_8H_{17} or $\text{C}_{14}\text{H}_{29}$. In another embodiment, R_H may contain a fluorocarbon group such as, for example, $\text{C}_n\text{F}_{2n+1}(\text{CH}_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24. R_F is independently selected and is a branched or straight chained fluorocarbon group (e.g., hydrocarbon/fluorocarbon group) (e.g., $\text{C}_n\text{F}_{2n+1}(\text{CH}_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24) wherein M^+ is a counteranion such as, for example, K^+ , Na^+ , or NH_4^+ . Preferred groups for R_H are, without limitation, $\text{C}_6\text{F}_{13}(\text{CH}_2)_2$ and $\text{C}_{10}\text{F}_{21}(\text{CH}_2)_2$ and preferred groups for R_F are, without limitation, C_4H_9 , C_8H_{17} , $\text{C}_{12}\text{H}_{25}$, and $\text{C}_{16}\text{H}_{33}$. X_I is a functional group such as hydroxy ($-\text{OH}$), alkyl (such that a phosphate bond is formed) and a phosphoamide groups (such as, for example, piperidine, morpholine, and the like). In one preferred embodiment, when M^+ is NH_4^+ , R_F and R_H are each independently selected and preferably range from $\text{C}_2\text{F}_5(\text{CH}_2)_2$ to $\text{C}_8\text{F}_{17}(\text{CH}_2)_2$.

Another preferred anionic surfactant according to the present invention is of the following formula:

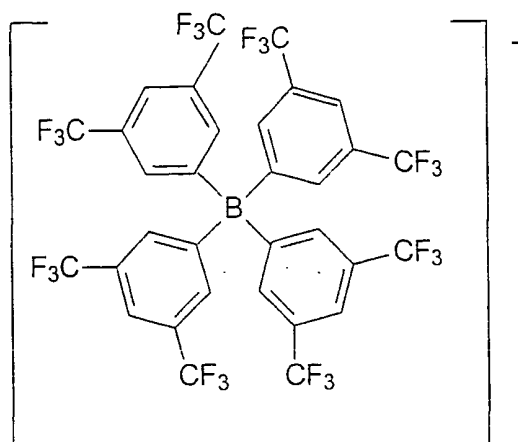


wherein R_F ranges from $\text{CF}_3(\text{CH}_2)_n$ to $\text{C}_{24}\text{F}_{49}(\text{CH}_2)_n$ wherein n ranges from 1 to 24..

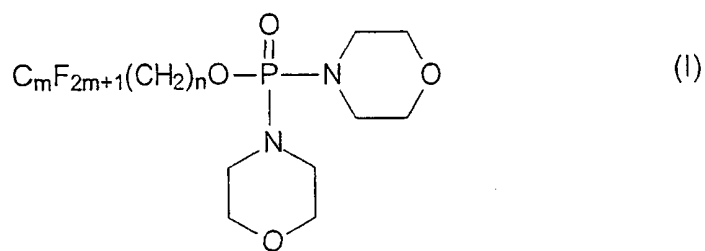
In other embodiments, the surfactant may be present in the form of a cationic surfactant. Examples of cationic surfactants are, without limitation, those represented by the formulas (VI) and (VII) (e.g., piperazinium phosphates):

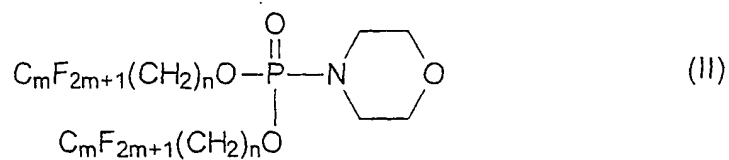


wherein R may be a branched or straight chained hydrocarbon group (e.g., $\text{C}_n\text{H}_{2n+1}$ wherein n ranges from 1 to 24) or a hydrocarbon/fluorocarbon group including, for example, $\text{C}_n\text{F}_{2n+1}(\text{CH}_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24, and X is a counteranion which may be a halogen (e.g., chloride, bromide), triflate, or "BarF" (anion of Kobayashi's reagent as set forth below):



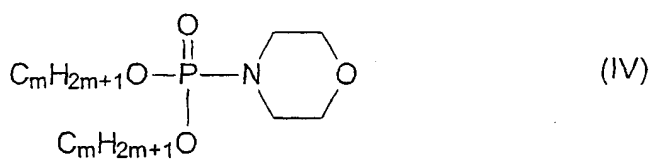
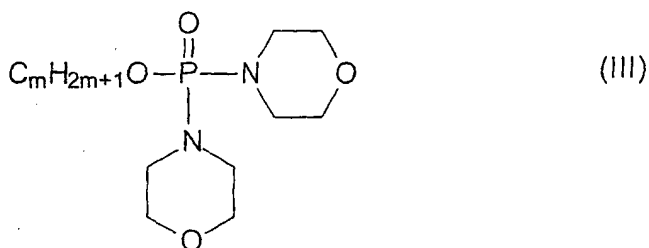
As an example, one or more morpholine units may be attached to the phosphate group. Embodiments represented by formulas (I) through (V) and (IX) and (X) describe such structures. Moreover, certain embodiments illustrate at least one fluorocarbon-containing group attached to the phosphate group. In one example, a fluorocarbon group is attached to a hydrocarbon spacer (e.g., 1 to 12 methylene units long) which is in turn attached to an oxygen atom and then the phosphorous atom as illustrated.





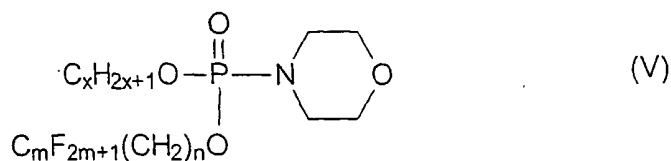
wherein m and n are each independently selected with m ranging from 1 to 24 and n ranging from 1 to 24. In a preferred embodiment, with respect to formula (I), m may be 6, 8, or 10, and n is 2. In a preferred embodiment, with respect to formula (II), m may be 6 or 8 and n is 2.

Hydrocarbon analogs of formulas (III) through (IV) are also encompassed.



wherein m is independently selected and ranges from 1 to 24. Preferably, m is 10.

Hybrid morpholine-containing surfactants can also be employed and are of the formula:



wherein x ranges from 1 to 24, m ranges from 1 to 24, and n ranges from 1 to 24. In a preferred embodiment, x is 8, m is 6, and n is 2.

The surfactants are believed to be potentially capable of reverse micelle and/or water-in carbon dioxide emulsion (i.e., microemulsion) formation. In certain embodiments, such compositions containing water may be present as homogeneous fluids or phases. Accordingly, in another aspect the invention provides a reverse micelle that comprises the composition of matter as defined herein and water. For the purposes of the invention, the term "reverse micelle" is defined as a micelle in which a water component is on the inner portion of the micelle. Thus, the "hydrophilic" segment of each surfactant present in the reverse micelle is on the inner portion of the micelle, while the "CO₂-philic" segment of each surfactant is on the outer portion of the micelle. As a result, the surfactants are useful in a number of applications. For example, the reverse micelles may be capable of encapsulating a number of pharmaceutically active agents therein, the selection of which is known to one skilled in the art. In other various embodiments, the surfactants can also be employed in cleaning or water removal in applications such as CO₂-based lithography, wherein a relatively small amount of water used in development can be dissolved and "swept away" into the phase containing predominantly carbon dioxide as well as synthesizing nanoparticles within microemulsion water pools. Organic (i.e. polymers) and inorganic nanoparticles can be also prepared using microemulsion water pools as reactors/templates. The surfactants can also be used in extraction applications, e.g., metal extractions from aqueous solutions.

Various amounts of water may be taken-up by the compositions of matter and methods of the present invention. For example, in one embodiment, the amount of water may range from above about 0, 5, 10, 20, 30, or 40 to about 60, 70, 80, 90, 95, 100, 150, or 200 percent based on the weight of the surfactant. More specifically, in accordance with the invention, the surfactant is capable of taking up 100 percent of its own weight in the form of water.

For the purposes of the invention, carbon dioxide is employed in a liquid or supercritical form. The composition of matter typically employs carbon dioxide as a continuous phase, with the composition of matter preferably comprising from about 50, 60, or 75 weight percent to about 80, 90, or 99 weight percent of carbon dioxide. If liquid CO₂ is used, the temperature employed during the process is preferably below 31°C. In one preferred embodiment, the CO₂ is utilized in a "supercritical" phase. As used herein, "supercritical" means that a fluid medium is at a temperature that is sufficiently high that it cannot be liquefied by pressure. The thermodynamic properties of CO₂ are reported in Hyatt, *J. Org. Chem.* **49**: 5097-5101 (1984); therein, it is stated that the critical temperature of CO₂ is about 31°C. In particular, the methods of the present invention are preferably carried out at a temperature range from about 20°C to about 60°C. The pressures employed preferably range from about 1000 psia (6.9 MPa) to about 5500 psia (37.9 MPa).

The composition of matter may also comprise components in addition to those described above. Preferably, these components do not react with the surfactant. Exemplary components include, but are not limited to, polymer modifier, water, rheology modifiers, plasticizing agents, antibacterial agents, flame retardants, and viscosity reduction modifiers. Co-solvents and co-surfactants may also be optionally employed. These components may be used if they do not react with the reactive functional polymer.

Exemplary co-solvents that could be used include, but are not limited to, alcohols (e.g., methanol, ethanol, and isopropanol); fluorinated and other halogenated solvents (e.g., chlorotrifluoromethane, trichlorofluoromethane, perfluoropropane, chlorodifluoromethane, and sulfur hexafluoride); amines (e.g.,

N-methyl pyrrolidone); amides (e.g., dimethyl acetamide); aromatic solvents (e.g., benzene, toluene, and xylenes); esters (e.g., ethyl acetate, dibasic esters, and lactate esters); ethers (e.g., diethyl ether, tetrahydrofuran, and glycol ethers); aliphatic hydrocarbons (e.g., methane, ethane, propane, butane, *n*-pentane, and hexanes); oxides (e.g., nitrous oxide); olefins (e.g., ethylene and propylene); natural hydrocarbons (e.g., isoprenes, terpenes, and *d*-limonene); ketones (e.g., acetone and methyl ethyl ketone); organosilicones; alkyl pyrrolidones (e.g., *N*-methyl pyrrolidone); paraffins (e.g., isoparaffin as well as other alkanes and paraffin waxes); petroleum-based solvents and solvent mixtures; and any other compatible solvent or mixture that is available and suitable. Mixtures of the above co-solvents may be used.

Exemplary co-surfactants that may possibly be used include, but are not limited to, longer chain alcohols (i.e., greater than or equal to C₈) such as octanol, decanol, dodecanol, cetyl alcohol, laurel alcohol, and the like; and species containing two or more alcohol groups or other hydrogen bonding functionalities; amides; amines; and other like components. Suitable other types of materials that are useful as co-surfactants are well known by those skilled in the art, and may be employed in the composition of matter of the present invention. Mixtures of the above may be used.

The invention also provides a method of separating the surfactant from the carbon dioxide and applying the surfactant to a substrate. In particular, the method comprises providing a composition of matter comprising at least one surfactant comprising (1) at least one phosphate group and (2) a solvent comprising carbon dioxide, wherein at least a portion of the surfactant is soluble in the solvent; and applying the composition of matter onto a substrate such that the carbon dioxide separates from the surfactant and wherein the surfactant coats the substrate. Techniques for separating and applying materials to a substrate are known in the art and are described, for example, in U.S. Patent No. 5,863,612 to DeSimone et al., the disclosure of which is incorporated herein by reference in its entirety, such as found on col. 5, line 47 through col. 6, line 11. Examples of methods for separating the surfactant include, without limitation,

boiling off the carbon dioxide and leaving the surfactant behind, and precipitation of the surfactant into a non-solvent either by introducing a non-solvent to a vessel or reactor containing the surfactant or the transfer of the vessel or reactor contents to another vessel containing a non-solvent for the surfactant. In one embodiment, the separation and application steps may be carried out together and include, as an example, passing (e.g., spraying or spray-drying) a solution containing the surfactant through an orifice to form particles, powder coatings, fibers, and other coatings on the substrates. A wide variety of substrates may be employed such as, without limitation, metals, organic polymers, inorganic polymers, textiles, and composites thereof. Exemplary substrates include, without limitation, integrated circuits, silicon wafers, silicon wafers with vias containing water, low-dielectric constant surfaces used as interlayer dielectrics on integrated circuits, a MEM, a porous material, a micro-porous material, a nano-porous material, a non-woven material, surfaces to be cleaned, surfaces to be treated with passivation layers, surfaces to be coated, surfaces to be treated with a self-assembled monolayer ("SAM"), photoresist coated surfaces, optical interfaces, optical relays, optical fibers, metallized surfaces, and micromirrors. These application techniques are demonstrated for liquid and supercritical solutions. The surfactant may form a low surface energy coating on the substrate. Examples of embodiments of substrates include, without limitation, textiles, papers, fiber optics, as well as other surfaces.

The compositions of matter, and in particular the surfactant, are useful in a number of applications such as, but not limited to, cleaning processes, solvent pool formation for polymerization processes, inorganic particle synthesis, and enzymatic reactions.

The composition of matter may include various amounts of surfactant. In a preferred embodiment, the composition of matter comprises from about 1, 2, 5, or 7 to about 5, 8, 10, 15, or 20 percent by weight of surfactant.

In another aspect, the invention provides a method of removing water from a composition of matter. The method comprises contacting a first composition of matter comprising water with a second composition of matter comprising: (1) at

least one surfactant comprising at least one phosphate group and (2) a solvent comprising carbon dioxide, wherein at least a portion of the surfactant is soluble in the solvent, such that the at least one surfactant removes at least a portion of the water from the first composition.

Embodiments describing the second composition of matter comprising the surfactant are set forth hereinabove, as well as for the solvent comprising carbon dioxide. In a preferred embodiment, the second composition of matter may be present as a homogeneous phase prior to and after contacting the first composition of matter, although other embodiments may also be contemplated, i.e., the second composition of matter may be heterogeneous.

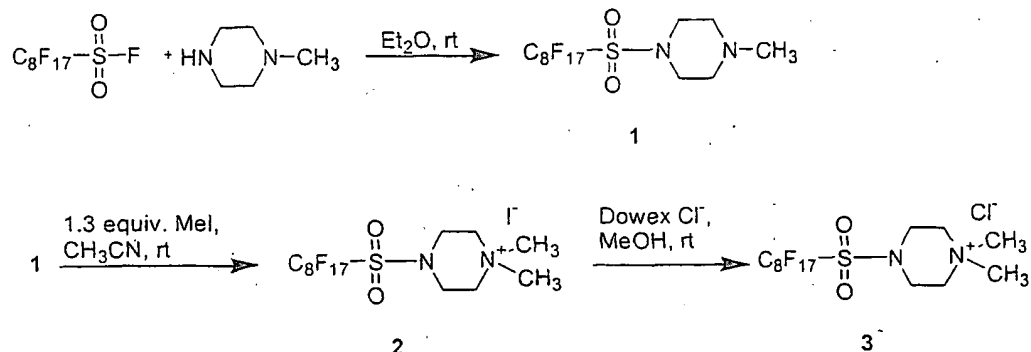
The first composition of matter comprising water may be present in the form of a number of embodiments, such as, for example, various articles of manufacture. Examples of such embodiments include, without limitation, integrated circuits, silicon wafers, silicon wafers with vias containing water, low-dielectric constant surfaces used as interlayer dielectrics on integrated circuits, a MEM, a porous material, a micro-porous material, a nano-porous material, a non-woven material, surfaces to be cleaned, surfaces to be treated with passivation layers, surfaces to be coated, surfaces to be treated with a self-assembled monolayer ("SAM"), photoresist coated surfaces, optical interfaces, optical relays, optical fibers, metallized surfaces, and micromirrors.

Methods for carrying out water removal may be conducted in systems, vessels, cells, and apparatuses known to one skilled in the art. Such systems, vessels, cells, and apparatuses include, without limitation, those capable of withstanding high pressure. Such equipment may be optionally contain agitation and heating means, the selection of which is known. The methods of water removal may be carried out by employing batch, continuous, and semi-continuous systems.

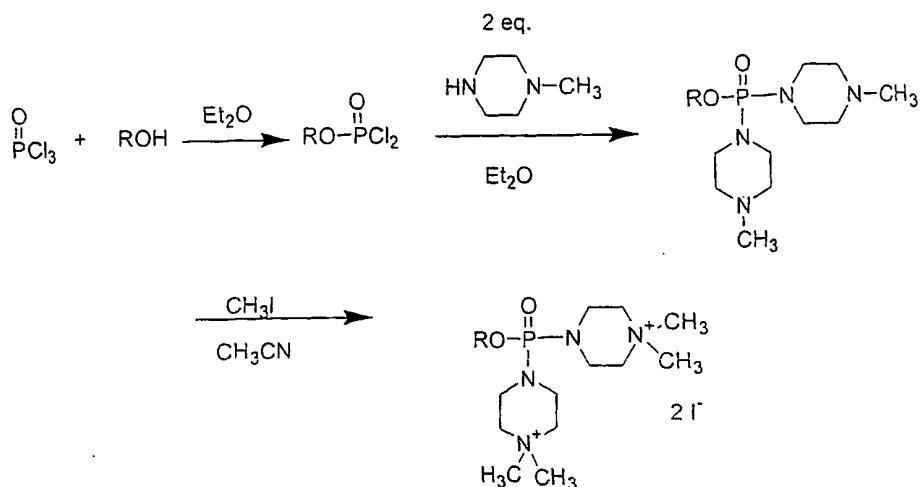
The surfactants that are employed in the invention may be formed by various techniques such as, for example, as set forth in Sadtler, V. M.; Jeanneaux, F.; Krafft, M. P.; Rábai, J.; Riess, J. G. *New Journal of Chemistry*. 1998, 609. Other synthesis routes may also be used. As an example,

morpholinolinosurfactants having fluorocarbon units attached thereto may be formed by first phosphorylation of a fluorinated alcohol by phosphorus oxychloride in the presence of $N(CH_2CH_3)_3$ and $(CH_3CH_2)_2O$. In one embodiment, the formation of di- and triesters and the chlorination of the alcohol can potentially be avoided by using dry ether and an excess of triethylamine, which results in the precipitation of triethylammonium chloride. (Perfluoroalkyl)alkyldimorpholinophosphates are subsequently obtained by a reaction with morpholine. The synthesis of bis[(F-alkyl)alkyl]monomorpholinophosphates also may involve the direct phosphorylation of F-alkylated alcohols by $OPCl_3$, followed by reaction with morpholine as described in Sadtler, V. M., et al.

Piperazine surfactants may be made by following the teachings of Katritzky, A.R., Davis, T.L., Rewcastle G.W., Rubel, G.O., and Pike, M.T., *Langmuir* 1988 4, 732-735. In accordance with the teachings set forth in Katritzky, et al., the following synthesis path may be followed for forming sulfonyl piperazinium compounds:

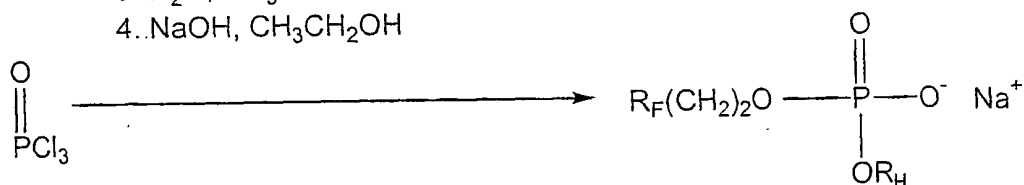


It is believed that one can form piperazinium phosphate surfactants by substituting *N*-methylpiperazine for morpholine in the synthetic pathway utilized for morpholinophosphate surfactant synthesis. The methylated nitrogen can then be quaternized by iodomethane to provide one variant of the desired compounds. An exemplary synthesis route is given below.



An embodiment illustrating a synthesis route for hybrid surfactants is illustrated according to the following scheme:

1. $R_F(CH_2)_2OH$, $N(CH_2CH_3)_3$, $(CH_3CH_2)_2O$
2. R_HOH , $N(CH_2CH_3)_3$, $(CH_3CH_2)_2O$
3. H_2O , CH_3CN
4. $NaOH$, CH_3CH_2OH



Neutral or anionic surfactants may be formed according to the above embodiment. It should be appreciated that these surfactants may be formed by other synthesis route. In the above synthesis route, R_H is C_mH_{2m+1} wherein m ranges from 1 to 24 and R_F may be C_nF_{2n+1} wherein n ranges from 1 to 24.

Various processing conditions (e.g., time and temperature) may be employed in the above method. In one embodiment, for example, steps 1 and 2 can take place at a temperature ranging from about 0°C to about 10°C with warming to room temperature for about 6 to about 12 hours under an inert

atmosphere (e.g., nitrogen or argon). Steps 3 and 4 take place under ambient conditions. The above synthesis employs equipment and techniques known in the art such as conventional glass round bottom flasks and stir bars. Other types of equipment can also be employed.

The following examples are intended to illustrate the present invention, and are not intended to limit the scope of the invention. In the examples, the morpholinophosphate surfactants were prepared as described in Sadtler, V. M. et al. These compounds were purified by silica gel chromatography. The anionic phosphate surfactants were synthesized by adapting known methods for preparing phosphates with fluorinated chains (see Krafft, M.-P.; Rolland, J.-P.; Vierling, P.; Riess, J. G. *New Journal of Chemistry* 1990, 14, 869.) and forming the sodium salt via neutralization in ethanol (see Romsted, L. S.; Zanette, D. *J. Phys. Chem.* 1988, 92, 4690.). The compounds exhibited the expected ^1H , ^{19}F , and ^{31}P NMR spectral patterns and also possessed adequate purity as determined by elemental analysis (Atlantic Microlabs, Norcross, GA), i.e., defined as generally within 0.3 % of the expected calculated analytical elemental percentages.)

Cloud point solubilities of surfactants and surfactants and water in carbon dioxide were carried out using a HIP variable volume pressure generator/view cell (maximum volume = 15 mL) containing a 0.5 inch thick sapphire window for viewing and a magnetic stir bar to agitate the solution. CO_2 was injected with the aid of an ISCO compression pump connected to the cell through high-pressure steel tubing. The cell was further attached to a Sensotec pressure transducer and an Omega thermocouple for pressure and temperature readouts, respectively. Measured amounts of surfactant and water were added at room temperature prior to pressurization with CO_2 . Samples were heated (controlled to $\pm 0.1^\circ\text{C}$) in the cell through the use of variac-controlled heating tape. Cloud points (judged as the reversible onset of a visually fully opaque solution) were taken on the cooling cycle by isothermally varying the pressure through volume changes facilitated by the hand-controlled piston. The cell was tipped at a

downward angle to aid in the observation of any phase-separated liquids. The cell was cleaned thoroughly between experiments.

UV-Vis spectra were acquired using a Perkin Elmer Lambda 40 spectrometer. Pressurized solutions were prepared in a 2.5 mL stainless steel cell, equipped with two 1 in. diameter \times 5/8 in. thick sapphire windows enclosing a 1 cm solution path length. Appropriate amount of surfactant and water were placed into the cell chamber, along with a 1/4 in. magnetic stir bar for agitation. A film of methyl orange (for a concentration of 5×10^{-5} M) was pre-cast and dried on one of the sapphire windows by addition of a stock solution *via* syringe. The chamber was tightly sealed, and the cell was pressurized and stirred until a clear, one-phase solution was present.

Examples 1-7

CO₂ Solubility Evaluation

The CO₂ solubility of the surfactants was carried out using a stainless steel view cell (2.5 mL internal volume) specially designed for high pressure studies. CO₂/surfactant solutions were held within two sapphire glass windows (sealed with Teflon® o-rings) which were, in turn, held in place by threaded steel caps fit to brass washers. Attached to the cell were steel tubings for injecting CO₂ and for venting the system, as well as a thermocouple and transducer to monitor internal temperature and pressure, respectively. CO₂ was transferred to the cell by way of an ISCO single pump compressor. A 1/4 inch magnetic bar was included to stir solutions. The chamber temperature could be raised with the aid of variac-controlled heating elements.

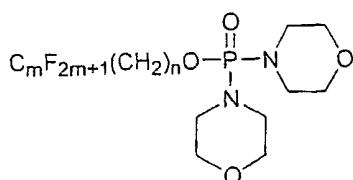
Experiments involved adding weighed amounts of the surfactants into the chamber along with the stir bar, tightly sealing the chamber, and injecting CO₂ to pressures between 1000 and 1500 psig at room temperature. The solutions were stirred and the temperatures increased at a rate of approximately 1°C/minute. Temperatures, pressures, and solution appearances were recorded. The results are listed in Table 1. The fluorosurfactants were soluble (i.e., dissolved in clear, transparent solutions) at concentrations of 10 percent (w/v)

over temperatures up to 60 °C, and exhibited no cloud points. Hydrocarbon analogs also proved to be soluble over similar pressure and temperature ranges.

Table 1. Solubilities of Morpholinophosphate Surfactants
in Dense CO₂ (2.5 mL volume)

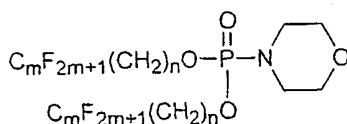
Surfactant	Weight of Sample (g)	Temperature Range (°C)	Pressure Range (psig)	Solubility
1	0.304	24-60	1070-2360	Clear Solution
2	0.298	24-53	1050-1820	Clear Solution
3	0.253	23-60	1100-2500	Clear Solution
4	0.259	25-59	1010-1760	Clear Solution
5	0.258	25-59	1070-2100	Clear Solution
6	0.253	23-59	1270-2960	Clear Solution
7	0.139	24-59	1250-2860	Clear Solution

Fluorosurfactants



Dimorpholinophosphate

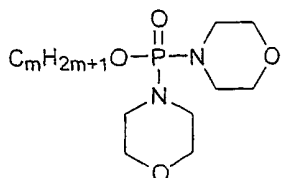
1: m=6, n=2
2: m=8, n=2
3: m=10, n=2



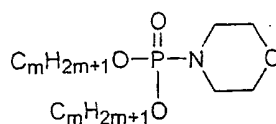
Monomorpholinophosphate

4: m=6, n=2
5: m=8, n=2

Hydrocarbon Analogs (control compounds)



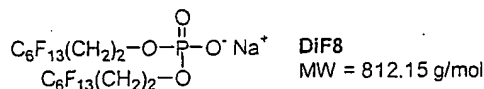
6: m=10



7: m=10

Example 8

Synthesis of Bis-[2-(*F*-hexyl)ethyl]phosphate, sodium salt



Bis-[2-(*F*-hexyl)ethyl]phosphate, sodium salt was synthesized according to the following example. Phosphorous oxychloride (5.27 g, 34.3 mmol) was added *via* syringe to 200 mL anhydrous diethyl ether, under nitrogen. The mixture was cooled to 0 °C, and a solution of 1*H*,1*H*,2*H*,2*H*-perfluorooctanol (25 g, 68.7 mmol) and triethylamine (17.4 g, 172 mmol) in 100 mL diethyl ether was slowly added, leading to the formation of white precipitate. The solution warmed to room temperature and was allowed to stir further under nitrogen overnight. The resulting white solid (triethylamine hydrochloride salt) was filtered and washed with 100 mL diethyl ether. The solvent and excess triethylamine was removed *via* rotary evaporation, providing an orange oil which was dissolved in 100 mL acetonitrile and 5 mL water. Two layers of immiscible liquids resulted, and the lower layer was isolated and dried under rotary evaporation. 22.5 g of crude as a viscous orange oil was provided. The oil was taken up in 100 mL ethanol. 2.28 g of NaOH (50 wt.% in water, 28.5 mmol NaOH) solution dissolved in 20 mL ethanol was slowly added. The solution precipitated into a gel, which was allowed to stir overnight. A white sticky solid (~ 3 g) was filtered and discarded, and the filtrate was concentrated *via* rotary evaporation and precipitated with diethyl ether. The resulting solid was dried and then taken up in 100 mL, and 1 g decolorizing carbon was added to the solution. This resulting solution was briefly stirred. The carbon was filtered off and solvent evaporated. 14.9 g (53 %) of the desired product after vacuum drying was provided (mp > 220 °C).

^1H NMR: (300 MHz: δ , CD_3OD) 4.13 (q, 4H; J_{HH} : 6.6 Hz, J_{HP} : 6.8 Hz; CH_2O), 2.55 (tt, 4H; J_{HH} : 6.6 Hz, J_{HF} : 19.2 Hz; CF_2CH_2)

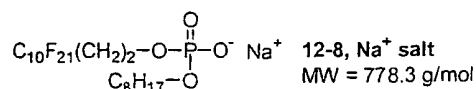
^{19}F NMR: (282 MHz: δ , CD_3OD) -83.0 (CF_2CF_3), -115.1 (CH_2CF_2), -123.2, -124.2, -125.0, (3 x CF_2), -127.7 (CF_2CF_3)

^{31}P NMR : (121 MHz: δ , CD_3OD) -0.65 ppm

Anal. Calcd. : C:23.66, H: 0.99; Found: C: 23.46; H 1.00.

Example 9

Synthesis of [2-(*F*-decyl)ethyl]octylphosphate, sodium salt



2-(*F*-decyl)ethyl]octylphosphate, sodium salt was synthesized as follows. Phosphorous oxychloride (1.3 g, 8.6 mmol) was added *via* syringe to 25 mL anhydrous diethyl ether, under argon. The mixture was cooled to 0 °C, and a solution of 1*H*,1*H*,2*H*,2*H*-perfluorododecanol (4.65 g, 8.2 mmol) and triethylamine (2.1 g, 20.8 mL) in 25 mL diethyl ether was slowly added. A white precipitate then formed. The mixture was stirred for 1 h at 0 °C, and a solution of 1-octanol (1.1 g, 8.2 mmol) and triethylamine (2.1 g, 20.8 mL) in 25 mL diethyl ether was added to the solution, resulting in the formation of more white precipitate. The solution was allowed to come to room temperature and was stirred under argon overnight. The white solid (triethylamine hydrochloride salt) was filtered and the filtrate was condensed *via* rotary evaporation and dissolved in a 20 mL mixture of acetonitrile and chloroform (19/1). 1 mL of water was slowly added and the solution was stirred overnight, providing a white solid, which was washed with acetonitrile and filtered. The solid was triturated in chloroform, the insoluble material filtered away, and the chloroform removed *via* rotary evaporation to provide 3.0 g of the neutral phosphate (48.4 percent) The corresponding sodium

salt was prepared in quantitative yield *via* neutralization with 1 equivalent of sodium hydroxide in ethanol (mp > 225 °C).

¹H NMR: (300 MHz: δ, CD₃OD) 4.14 (q, 2H; *R_F* chain CH₂O), 3.84 (q, 2H; *R_H* chain CH₂O), 2.55 (m, 2H; CF₂CH₂), 1.6 (m, 2H; CH₂CH₂CH₂O), 1.2-1.4 (m, 10H), 0.88 (t, 3H; *J_{HH}*: 7.1 Hz)

¹⁹F NMR: (282 MHz: δ, CD₃OD) -81.4 (CF₂CF₃), -113.3 (CH₂CF₂), -121.4, -122.4, -123.4, (7 x CF₂), -126.0 (CF₂CF₃)

³¹P NMR: (121 MHz: δ, CD₃OD) 1.53

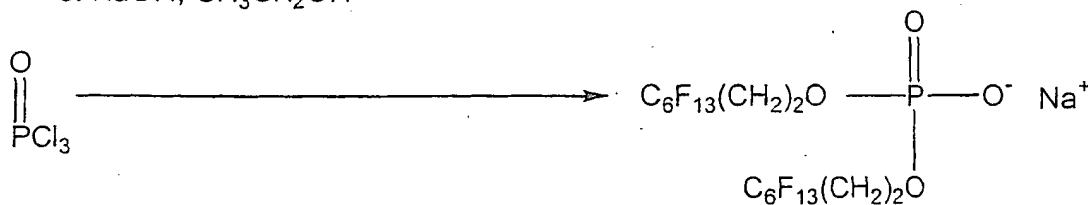
Anal. Calcd.: C: 30.86, H: 2.72; Found: C: 30.73, H: 2.62.

Example 10

Synthesis of Di-Fluoro-Chained Analog Surfactant

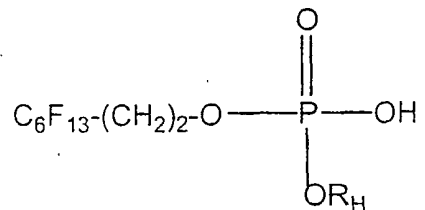
Di-fluoro chained analog surfactants were formed according to the following synthesis route:

1. 2 eq. C₆F₁₃(CH₂)₂OH, N(CH₂CH₃)₂, (CH₃CH₂)₂O
2. H₂O, CH₃CN
3. NaOH, CH₃CH₂OH



Example 11**Cloud Point Evaluation**

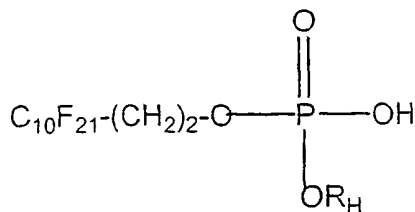
The solubilities of various surfactants of the formula described below were evaluated in carbon dioxide:



wherein R_H is C_4H_9 ("8-4"), C_8H_{17} ("8-8"), $\text{C}_{12}\text{H}_{25}$ ("8-12"), and $\text{C}_{16}\text{H}_{33}$ ("8-16"). Cloud point curves for these surfactants are set forth in FIG 2.

Example 12**Cloud Point Evaluation**

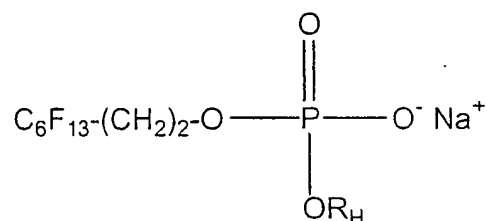
The solubilities of various surfactants of the formula described below were evaluated in carbon dioxide:



wherein R_H is C_4H_9 ("12-4"), C_8H_{17} ("12-8"), and $\text{C}_{12}\text{H}_{25}$ ("12-12"). Cloud point curves for these surfactants are set forth in FIG 3.

Example 13**Cloud Point Evaluation (Anionic Surfactant)**

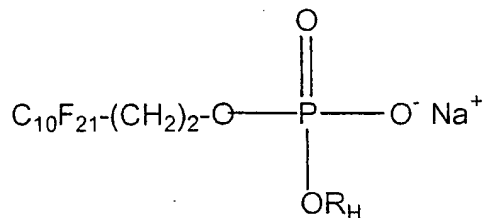
The solubility of an anionic surfactant of the formula described below was evaluated in carbon dioxide:



wherein R_H is C_4H_9 ("8-4"). A cloud point curve for this surfactant is set forth in FIG. 4.

Example 14**Cloud Point Evaluation (Anionic Surfactants)**

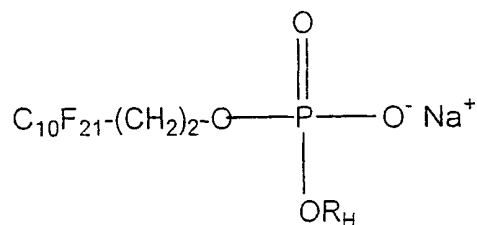
The solubility of an anionic surfactant of the formula described below was evaluated in carbon dioxide:



wherein R_H is C_4H_9 ("12-4") and C_8H_{17} ("12-8"). A cloud point curve for this surfactant is set forth in FIG. 5.

Example 15**Water Uptake Evaluation (Anionic Surfactant)**

The water uptake capability of an anionic surfactant was evaluated wherein the surfactant is of the formula:



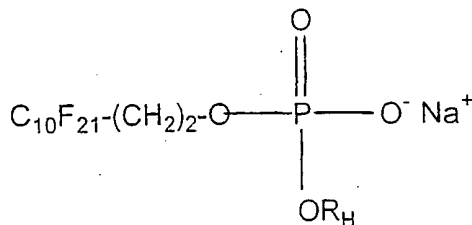
wherein R_H is C_4H_9 ("12-4").

The concentration of surfactant in carbon dioxide was 2.5 weight percent. The results are set forth in FIG. 6.

Example 16

Water Uptake Evaluation (Anionic Surfactants)

The water uptake capability of an anionic surfactant was evaluated wherein the surfactant is of the formula:

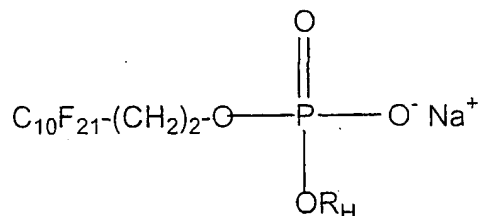


wherein R_H is C_4H_{17} ("12-8").

The concentration of surfactant in carbon dioxide was 2.5 weight percent. The results are set forth in FIG. 7.

Example 17**Water Uptake Evaluation (Anionic Surfactants)**

The water uptake capability of an anionic surfactant was evaluated wherein the surfactant is of the formula:

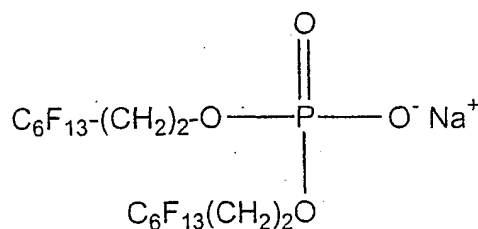


wherein R_H is $\text{C}_{12}\text{H}_{25}$ ("12-12").

The concentration of surfactant in carbon dioxide was 2.5 weight percent.
The results are set forth in FIG. 8.

Example 18**Water Uptake Evaluation (Anionic Surfactants)**

The water uptake capability of an anionic surfactant was evaluated wherein the surfactant is of the formula:



The concentration of surfactant in carbon dioxide was 2.5 weight percent.
The results are set forth in FIG. 9.

In the specification, drawings, and examples there have been disclosed typical preferred embodiments of the invention and, although specific terms are employed, they are used in a generic and descriptive sense only and not for the purposes of limitation, the scope of the invention being set forth in the following claims.

THAT WHICH IS CLAIMED:

1. A method of removing water from a composition of matter, said method comprising:

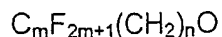
contacting a first composition of matter comprising water with a second composition of matter comprising: (1) at least one surfactant comprising at least one phosphate group and (2) a solvent comprising carbon dioxide, wherein at least a portion of the surfactant is soluble in the solvent, such that the at least one surfactant removes at least a portion of the water from the first composition

2. The method according to Claim 1, wherein the first composition of matter is present in an embodiment selected from the group consisting of integrated circuits, silicon wafers, silicon wafers with vias containing water, low-dielectric constant surfaces used as interlayer dielectrics on integrated circuits, a MEM, a porous material, a micro-porous material, a nano-porous material, a non-woven material, surfaces to be cleaned, surfaces to be treated with passivation layers, surfaces to be coated, surfaces to be treated with a self-assembled monolayer ("SAM"), photoresist coated surfaces, optical interfaces, optical relays, optical fibers, metallized surfaces, and micromirrors.

3. The method according to Claim 1, wherein the at least one surfactant comprises at least one fluorocarbon group.

4. The method according to Claim 3, wherein the at least one fluorocarbon group comprises a hydrocarbon spacer attached to an oxygen atom, wherein the oxygen atom is attached to the phosphate group.

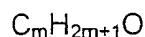
5. The method according to Claim 3, wherein the at least one fluorocarbon group is of the formula:



wherein m ranges from 1 to 24 and n ranges from 1 to 24.

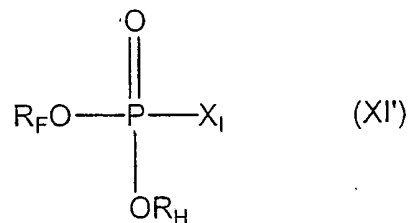
6. The method according to Claim 5, wherein said surfactant further comprises a hydrocarbon-containing group attached to the phosphate group.

7. The method according to Claim 6, wherein the hydrocarbon-containing chain is of the formula:



wherein m ranges from 1 to 24.

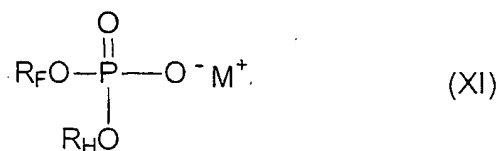
8. The method according to Claim 1, wherein the surfactant is represented by the formula:



wherein R_H is, C_nH_{2n+1} wherein n ranges from 1 to 24 or $C_nF_{2n+1}(CH_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24 or C_pH_{2p+1} wherein p ranges from 1 to 24, R_F is $C_nF_{2n+1}(CH_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24, and X_I is a functional group selected from the group consisting of hydroxy, alkyl, and phosphoamide groups.

9. The method according to Claim 1, wherein the surfactant is an anionic surfactant.

10. The method according to Claim 9, wherein the anionic surfactant is represented by the formula:

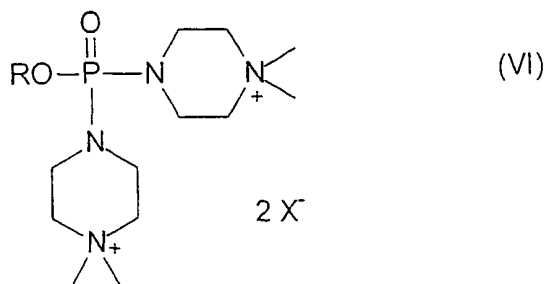


wherein R_H is, $\text{C}_n\text{H}_{2n+1}$ wherein n ranges from 1 to 24 or $\text{C}_n\text{F}_{2n+1}(\text{CH}_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24, R_F is $\text{C}_n\text{F}_{2n+1}(\text{CH}_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24, and M is K^+ , Na^+ , or NH_4^+ .

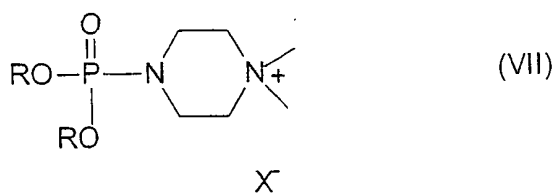
11. The method according to Claim 9, wherein R_F is selected from the group consisting of $\text{C}_6\text{F}_{13}(\text{CH}_2)_2$ and $\text{C}_{10}\text{F}_{21}(\text{CH}_2)_2$, R_H is selected from the group consisting of C_4H_9 , C_8H_{17} , $\text{C}_{12}\text{H}_{25}$, and $\text{C}_{16}\text{H}_{33}$, and M is Na^+ .

12. The method according to Claim 1, wherein the surfactant is a cationic surfactant.

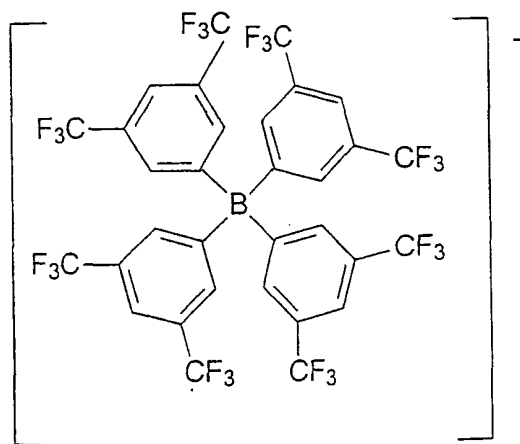
13. The method according to Claim 12, wherein the cationic surfactant is selected from the group consisting of:



and



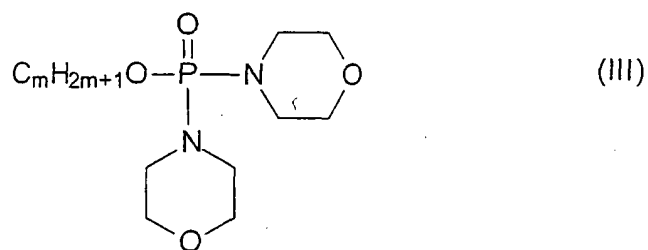
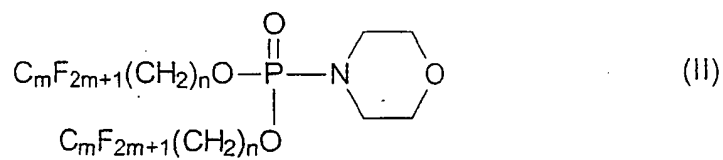
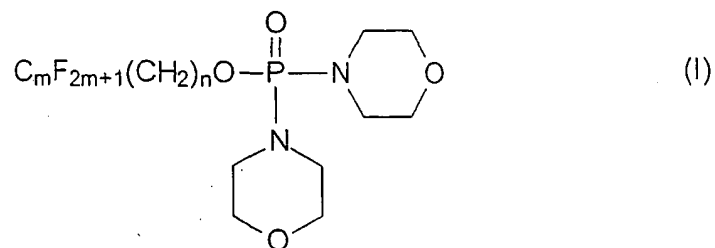
wherein R is selected from the group consisting of C_nH_{2n+1} wherein n ranges from 1 to 24 and $C_nF_{2n+1}(CH_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24, and wherein X is a counteranion selected from the group consisting of halogen, triflate, and

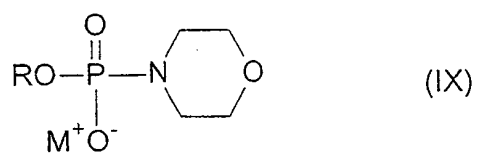
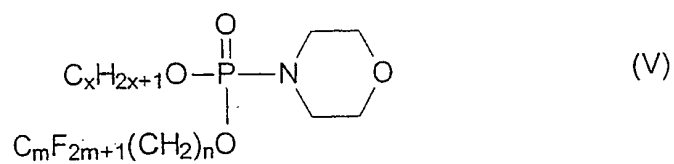
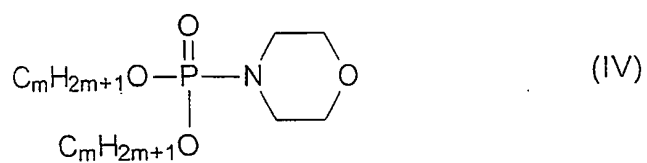


14 The method according to Claim 1, wherein the

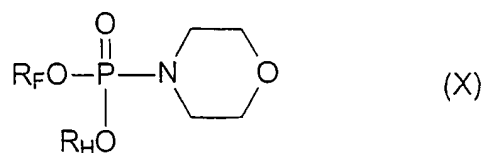
surfactant comprises one or more one or more morpholine units attached to the phosphate group.

15. The method according to Claim 14, wherein the surfactant is selected from the group consisting of





and



wherein m and n are independently selected with m ranging from 1 to 24 and n ranging from 1 to 24, x ranges from 1 to 24, M is selected from the group consisting of K^+ , Na^+ , and NH_4^+ , R is $\text{C}_p\text{H}_{2p+1}$ wherein p ranges from 1 to 24 R_F is $\text{C}_n\text{F}_{2n+1}$ wherein n ranges from 1 to 24, and R_H is either: (1) $\text{C}_n\text{H}_{2n+1}$ wherein n ranges from 1 to 24 or (2) $\text{C}_n\text{F}_{2n+1}(\text{CH}_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24.

16. The method according to Claim 1, wherein the carbon dioxide is supercritical carbon dioxide.

17. The method according to Claim 1, wherein the carbon dioxide is liquid carbon dioxide.

18. The method according to Claim 1, wherein said surfactant is present in said composition of matter in an amount ranging from about 1 to about 10 percent based on the weight of the surfactant.

19. The method according to Claim 1, further comprising at least one additional component selected from the group consisting of polymer modifier, rheology modifiers, plasticizing agents, antibacterial agents, flame retardants, viscosity reduction modifiers, co-solvents, and co-surfactants.

20. The method according to Claim 1, wherein the water and the surfactant in the second composition of matter form a reverse micelle.

21. The method according to Claim 1, wherein the water is

present in the second composition of matter in an amount ranging from above about 0 to about 200 percent based on the weight of the surfactant.

22. A method of applying a surfactant to a substrate, said method comprising:

providing a composition of matter comprising at least one surfactant comprising (1) at least one phosphate group and (2) a solvent comprising carbon dioxide, wherein at least a portion of the surfactant is soluble in the solvent; and

applying the composition of matter onto a substrate such that the carbon dioxide separates from the surfactant and wherein the surfactant coats the substrate.

23. The method according to Claim 22, wherein the substrate is selected from the group consisting of metals, organic polymers, inorganic polymers, textiles, and composites thereof

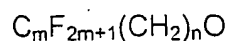
24. The method according to Claim 23, wherein the substrate is selected from the group consisting of integrated circuits, silicon wafers, silicon wafers with vias containing water, low-dielectric constant surfaces used as interlayer dielectrics on integrated circuits, a MEM, a porous material, a micro-porous material, a nano-porous material, a non-woven material, surfaces to be cleaned, surfaces to be treated with passivation layers, surfaces to be coated, surfaces to be treated with a self-assembled monolayer, photoresist coated surfaces, optical interfaces, optical relays, optical fibers, metallized surfaces, and micromirrors.

25. The method according to Claim 22, wherein the at least one surfactant comprises at least one fluorocarbon group.

26. The method according to Claim 25, wherein the at

least one fluorocarbon group comprises a hydrocarbon spacer attached to an oxygen atom, wherein the oxygen atom is attached to the phosphate group.

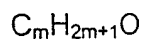
27. The method according to Claim 26, wherein the at least one fluorocarbon group is of the formula:



wherein m ranges from 1 to 24 and n ranges from 1 to 24.

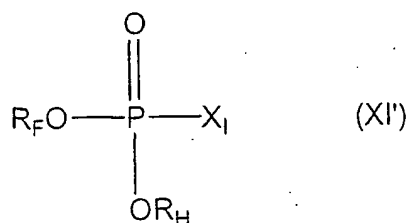
28. The method according to Claim 26, wherein said surfactant further comprises a hydrocarbon-containing group attached to the phosphate group.

29. The method according to Claim 28, wherein the hydrocarbon-containing chain is of the formula:



wherein m ranges from 1 to 24.

30. The method according to Claim 22, wherein the surfactant is represented by the formula:

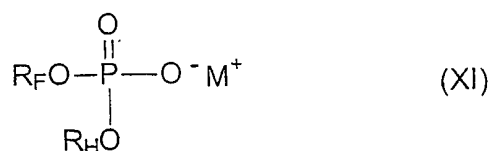


wherein R_H is, C_nH_{2n+1} wherein n ranges from 1 to 24 or $C_nF_{2n+1}(CH_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24 or C_pH_{2p+1} wherein p ranges from 1

to 24, R_F is $C_nF_{2n+1}(CH_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24, and X_1 is a functional group selected from the group consisting of hydroxy, alkyl, and phosphoamide groups.

31. The method according to Claim 22, wherein the surfactant is an anionic surfactant.

32. The method according to Claim 30, wherein the anionic surfactant is represented by the formula:

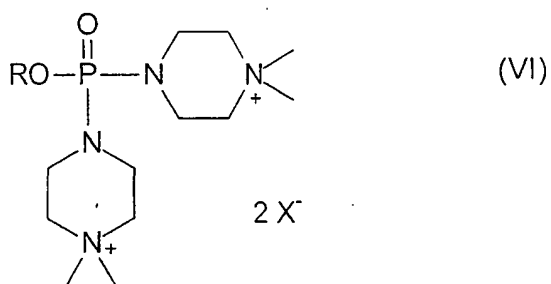


wherein R_H is, C_nH_{2n+1} wherein n ranges from 1 to 24 or $C_nF_{2n+1}(CH_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24 R_F is $C_nF_{2n+1}(CH_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24, and M is K^+ , Na^+ , or NH_4^+ .

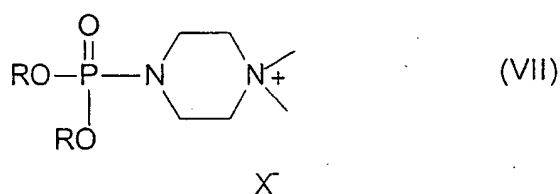
33. The method according to Claim 32, wherein R_F is selected from the group consisting of $C_6F_{13}(CH_2)_2$ and $C_{10}F_{21}(CH_2)_2$, R_H is selected from the group consisting of C_4H_9 , C_8H_{17} , $C_{12}H_{25}$, and $C_{16}H_{33}$, and M is Na^+ .

34. The method according to Claim 22, wherein the surfactant is a cationic surfactant.

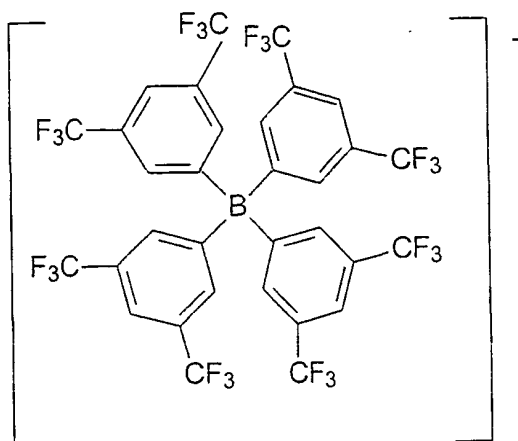
35. The method according to Claim 34, wherein the cationic surfactant is selected from the group consisting of:



and

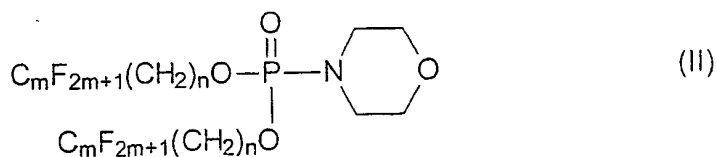
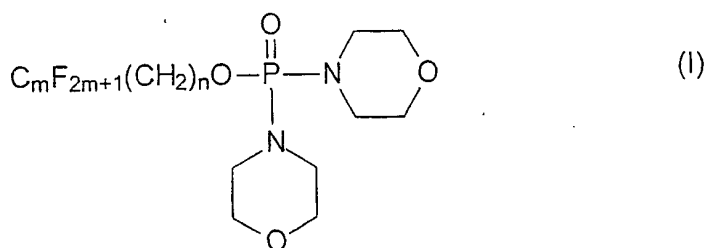


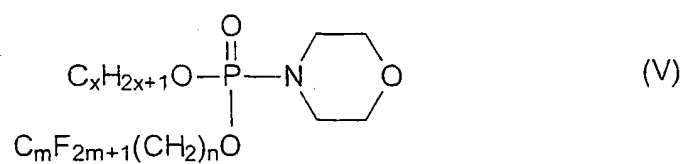
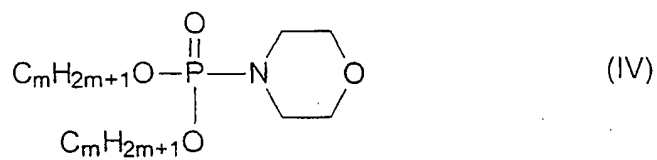
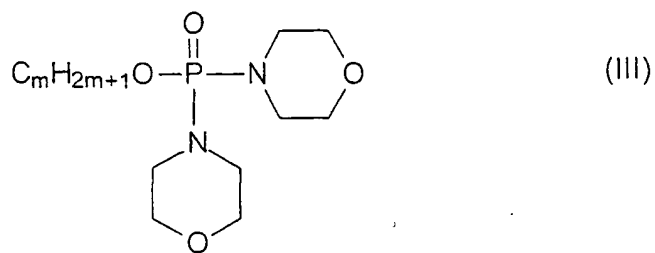
wherein R is selected from the group consisting of C_nH_{2n+1} wherein n ranges from 1 to 24 and $C_nF_{2n+1}(CH_2)_m$ wherein m ranges from 1 to 24, and wherein X is a counteranion selected from the group consisting of halogen, triflate, and

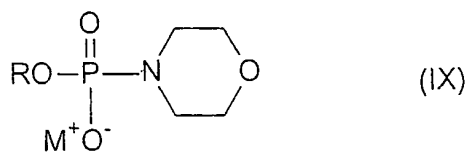


36. The method according to Claim 22; wherein the surfactant comprises one or more one or more morpholine units attached to the phosphate group.

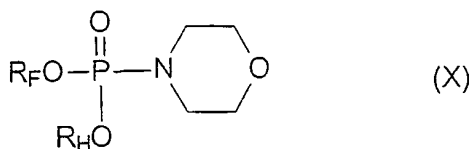
37. The method according to Claim 36, wherein the surfactant is selected from the group consisting of







and



wherein m and n are independently selected with m ranging from 1 to 24 and n ranging from 1 to 24, x ranges from 1 to 24, M is selected from the group consisting of K^+ , Na^+ , and NH_4^+ , R is $\text{C}_p\text{H}_{2p+1}$ wherein p ranges from 1 to 24 R_F is $\text{C}_n\text{F}_{2n+1}$ wherein n ranges from 1 to 24, and R_H is either: (1) $\text{C}_n\text{H}_{2n+1}$ wherein n ranges from 1 to 24 or (2) $\text{C}_n\text{F}_{2n+1}(\text{CH}_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24.

38. The method according to Claim 22, wherein the carbon dioxide is supercritical carbon dioxide.

39. The method according to Claim 22, wherein the carbon dioxide is liquid carbon dioxide.

40. The method according to Claim 22, wherein said surfactant is present in said composition of matter in an amount ranging from about 1 to about 10 percent based on the weight of the surfactant.

41. The method according to Claim 22, further comprising

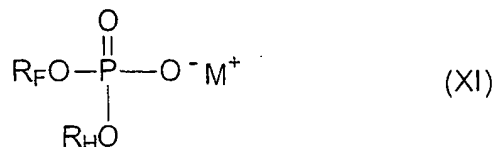
at least one additional component selected from the group consisting of polymer modifier, rheology modifiers, plasticizing agents, antibacterial agents, flame retardants, viscosity reduction modifiers, co-solvents, and co-surfactants.

42. The method according to Claim 22, wherein the composition of matter comprises water.

43. A composition of matter comprising:

- (1) at least one anionic surfactant comprising at least one phosphate group; and
- (2) a solvent comprising carbon dioxide, wherein at least a portion of the surfactant is soluble in the solvent.

wherein the anionic surfactant is represented by the formula:



wherein R_H is, $\text{C}_n\text{H}_{2n+1}$ wherein n ranges from 1 to 24 or $\text{C}_n\text{F}_{2n+1}(\text{CH}_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24, R_F is $\text{C}_n\text{F}_{2n+1}(\text{CH}_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24, and M is K^+ or Na^+ .

44. The composition of matter according to Claim 43, wherein R_F is selected from the group consisting of $\text{C}_6\text{F}_{13}(\text{CH}_2)_2$ and $\text{C}_{10}\text{F}_{21}(\text{CH}_2)_2$, R_H is selected from the group consisting of C_4H_9 , C_8H_{17} , $\text{C}_{12}\text{H}_{25}$, and $\text{C}_{16}\text{H}_{33}$, and M is Na^+ .

45. The composition of matter according to Claim 43, wherein the

carbon dioxide is supercritical carbon dioxide.

46. The composition of matter according to Claim 43, wherein the carbon dioxide is liquid carbon dioxide.

47. The composition of matter according to Claim 43, wherein said surfactant is present in said composition of matter in an amount ranging from about 1 to about 10 percent based on the weight of the surfactant.

48. The composition of matter according to Claim 43, further comprising at least one additional component selected from the group consisting of polymer modifier, rheology modifiers, plasticizing agents, antibacterial agents, flame retardants, viscosity reduction modifiers, co-solvents, and co-surfactants.

49. The composition of matter according to Claim 43, wherein the composition of matter comprises water.

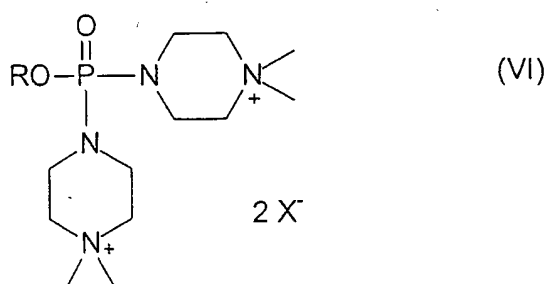
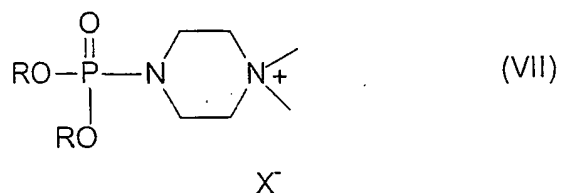
50. A reverse micelle comprising the composition of matter as defined by Claim 49.

51. The reverse micelle according to Claim 50, wherein the water is present in an amount ranging from above about 0 to about 200 percent based on the weight of the surfactant.

52. A composition of matter comprising:

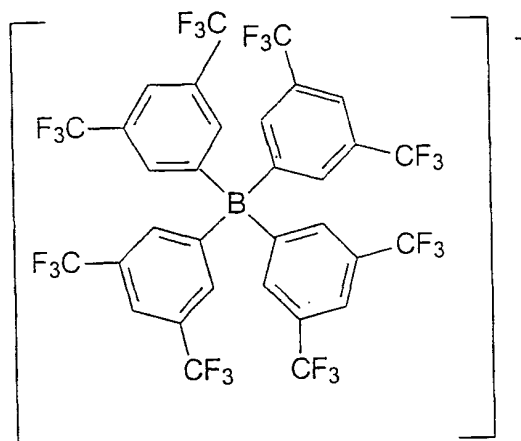
- (1) at least one cationic surfactant comprising at least one phosphate group; and
- (2) a solvent comprising carbon dioxide, wherein at least a portion of the surfactant is soluble in the solvent.

53. The composition of matter according to Claim 52, wherein the cationic surfactant is selected from the group consisting of:



and

wherein R is selected from the group consisting of C_nH_{2n+1} wherein n ranges from 1 to 24 and $C_nF_{2n+1}(CH_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24, and wherein X is a counteranion selected from the group consisting of halogen, triflate, and



54. The composition of matter according to Claim 52, wherein the carbon dioxide is supercritical carbon dioxide.

55. The composition of matter according to Claim 52, wherein the carbon dioxide is liquid carbon dioxide.

56. The composition of matter according to Claim 52, wherein said surfactant is present in said composition of matter in an amount ranging from about 1 to about 10 percent based on the weight of the surfactant.

57. The composition of matter according to Claim 52, further comprising at least one additional component selected from the group consisting of polymer modifier, rheology modifiers, plasticizing agents, antibacterial agents, flame retardants, viscosity reduction modifiers, co-solvents, and co-surfactants.

58. The composition of matter according to Claim 52, wherein the composition of matter comprises water.

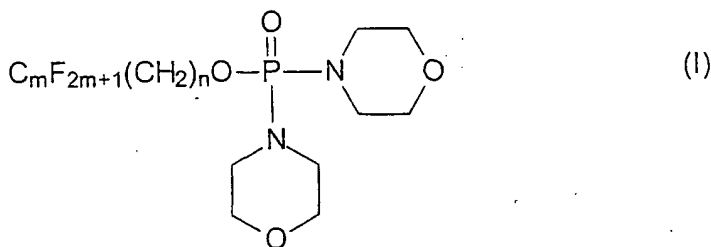
59. A reverse micelle comprising the composition of matter as defined by Claim 58.

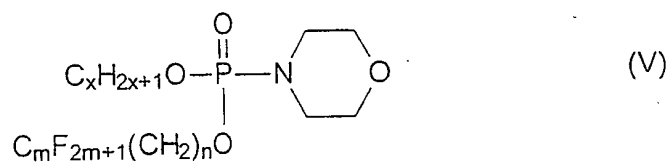
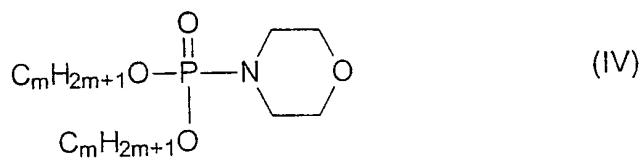
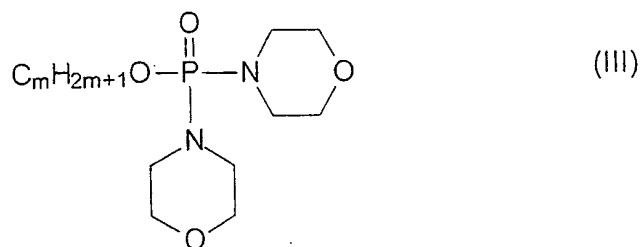
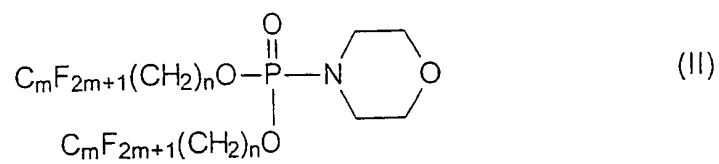
60. The reverse micelle according to Claim 59, wherein the water is present in an amount ranging from above about 0 to about 200 percent based on the weight of the surfactant.

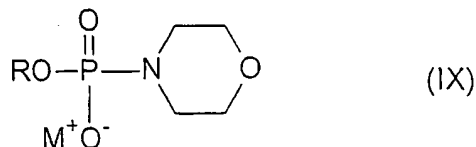
61. A composition of matter comprising:

- (1) at least one surfactant comprising at least one phosphate group; and
 - (2) a solvent comprising carbon dioxide, wherein at least a portion of the surfactant is soluble in the solvent;
- wherein the at least one surfactant comprises one or more one or more morpholine units attached to the phosphate group.

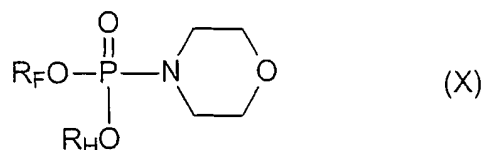
62. The composition of matter according to Claim 61, wherein the surfactant is selected from the group consisting of







and



wherein m and n are independently selected with m ranging from 1 to 24 and n ranging from 1 to 24, x ranges from 1 to 24, M is selected from the group consisting of K^+ , Na^+ , and NH_4^+ , R is $\text{C}_p\text{H}_{2p+1}$ wherein p ranges from 1 to 24 R_F is $\text{C}_n\text{F}_{2n+1}$ wherein n ranges from 1 to 24, and R_H is either: (1) $\text{C}_n\text{H}_{2n+1}$ wherein n ranges from 1 to 24 or (2) $\text{C}_n\text{F}_{2n+1}(\text{CH}_2)_m$ wherein n ranges from 1 to 24 and m ranges from 1 to 24.

63. The composition of matter according to Claim 61, wherein the carbon dioxide is supercritical carbon dioxide.

64. The composition of matter according to Claim 61, wherein the carbon dioxide is liquid carbon dioxide.

65. The composition of matter according to Claim 61, wherein said surfactant is present in said composition of matter in an amount ranging from about 1 to about 10 percent based on the weight of the surfactant.

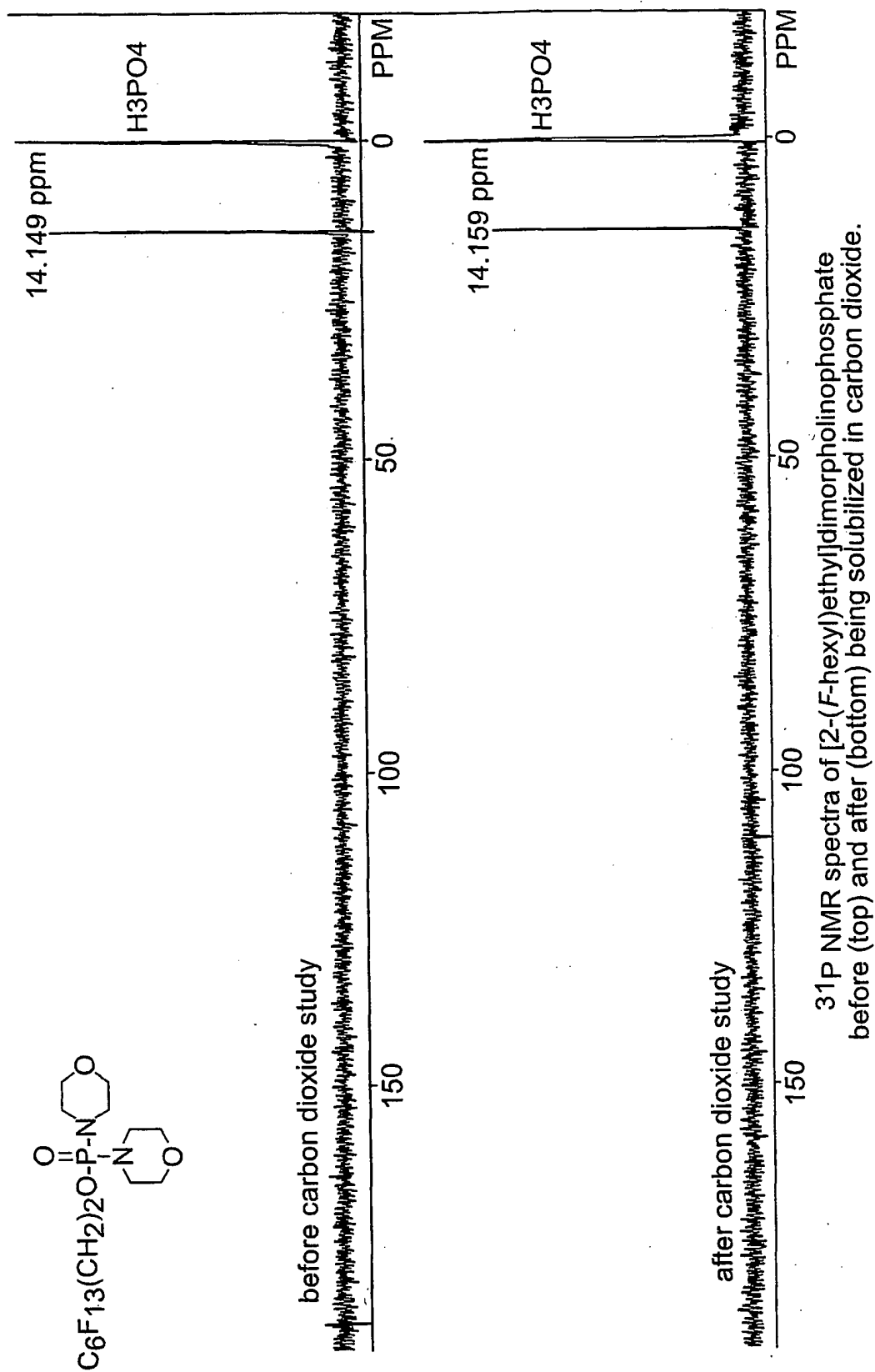
66. The composition of matter according to Claim 61, further comprising at least one additional component selected from the group consisting of polymer modifier, rheology modifiers, plasticizing agents, antibacterial agents, flame retardants, viscosity reduction modifiers, co-solvents, and co-surfactants.

67. The composition of matter according to Claim 61, wherein the composition of matter comprises water.

68. A reverse micelle comprising the composition of matter as defined by Claim 67.

69. The reverse micelle according to Claim 68, wherein the water is present in an amount ranging from above about 0 to about 200 percent based on the weight of the surfactant.

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**FIG. 1.**

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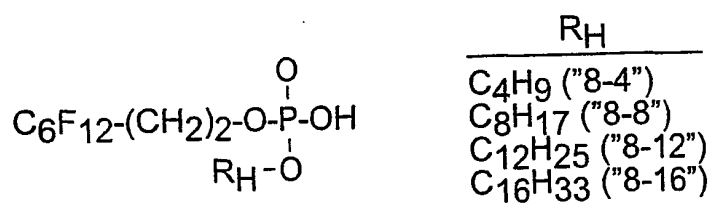
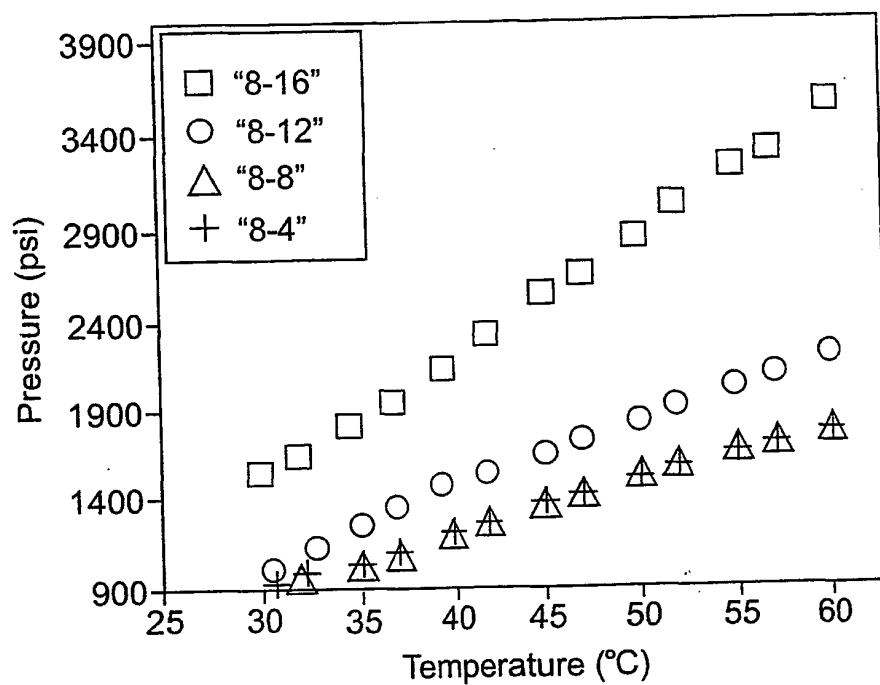


FIG. 2.

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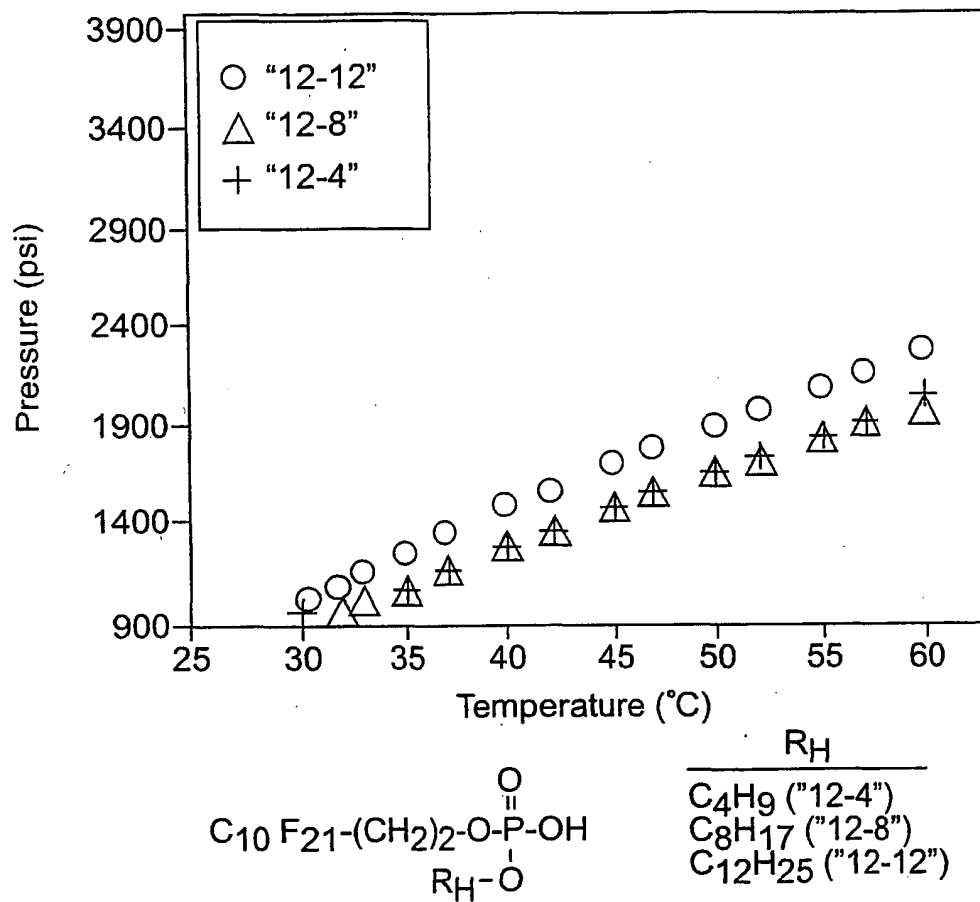


FIG. 3.

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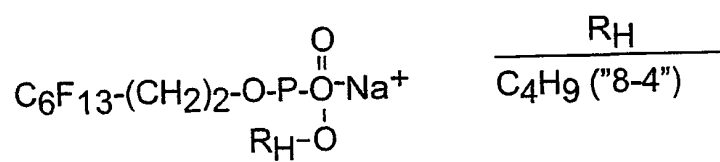
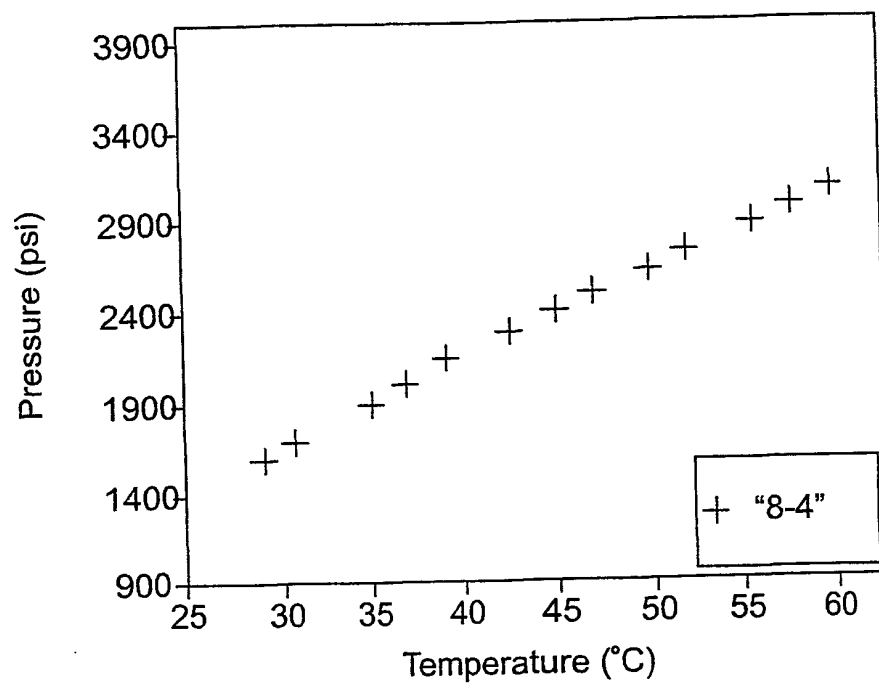


FIG. 4.

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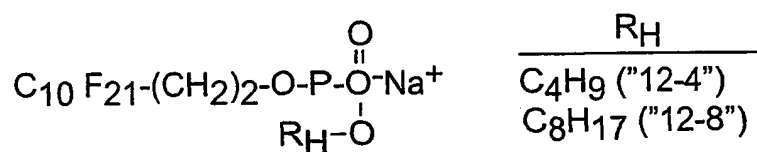
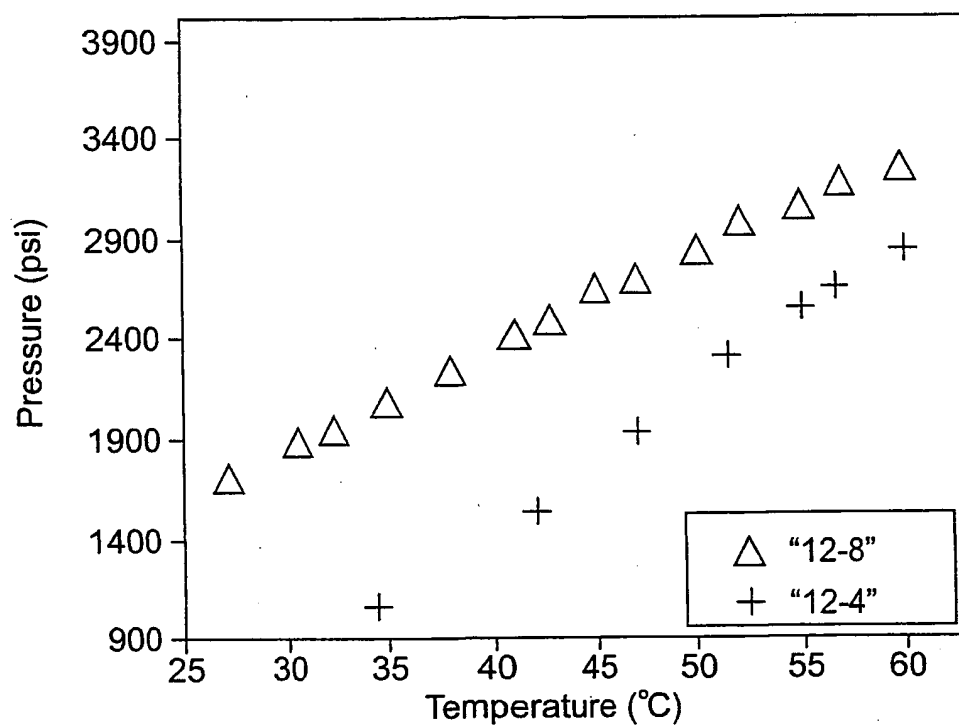
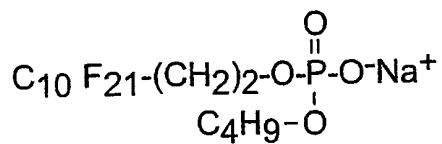
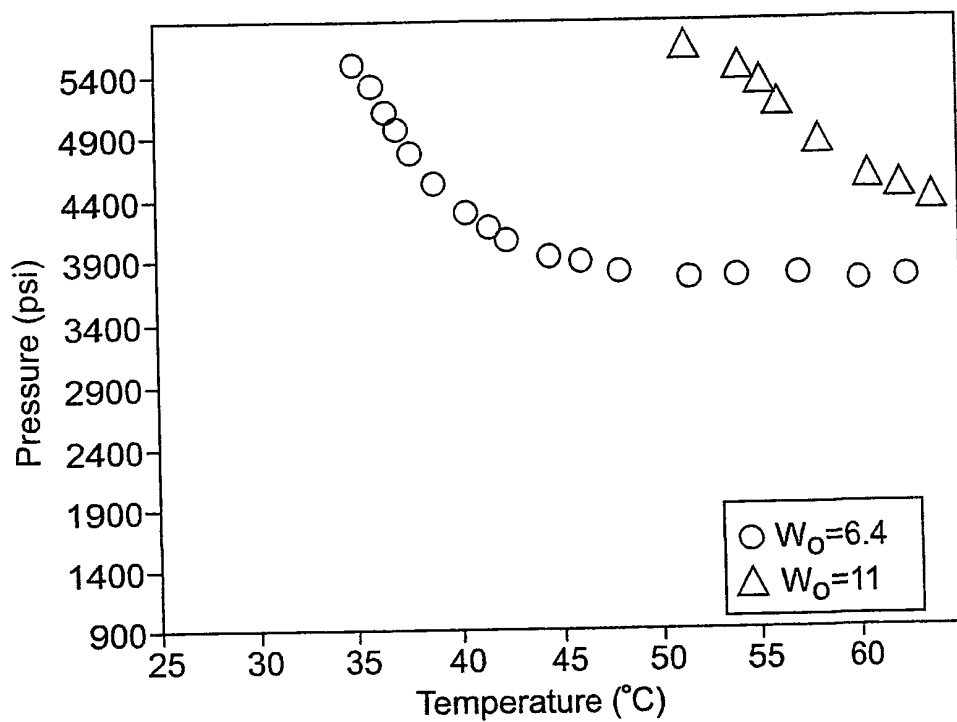


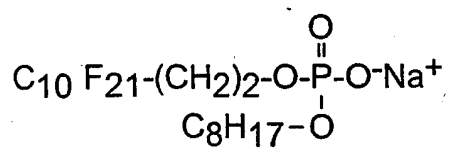
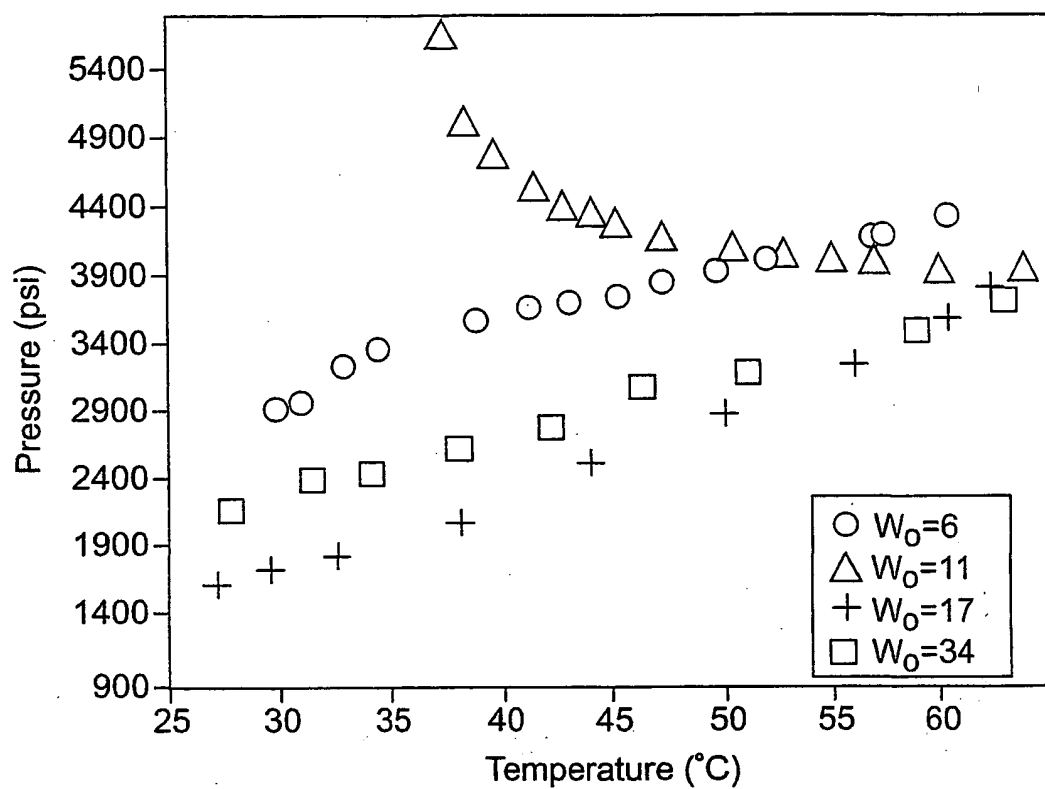
FIG. 5.

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2.5 wt.% in CO₂

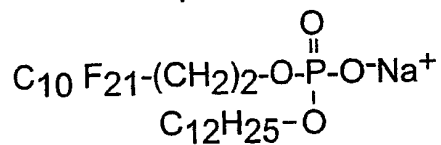
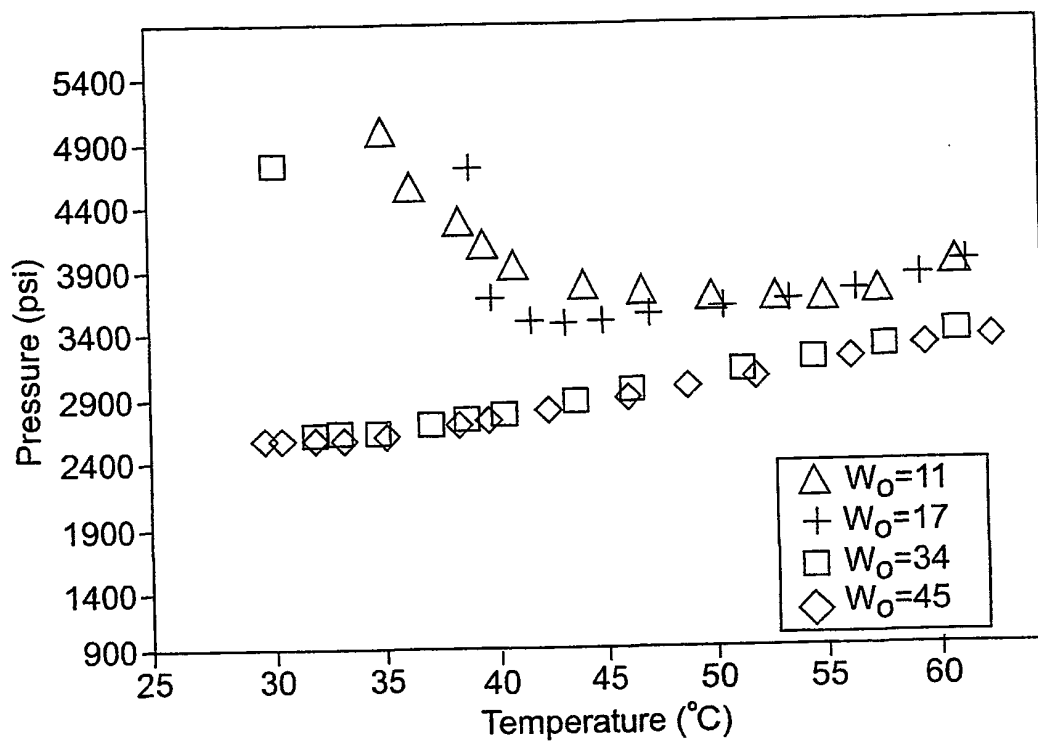
$$W_0 = \frac{\text{moles H}_2\text{O}}{\text{moles surf.}}$$
FIG. 6.

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2.5 wt.% in CO₂

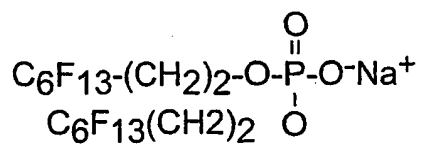
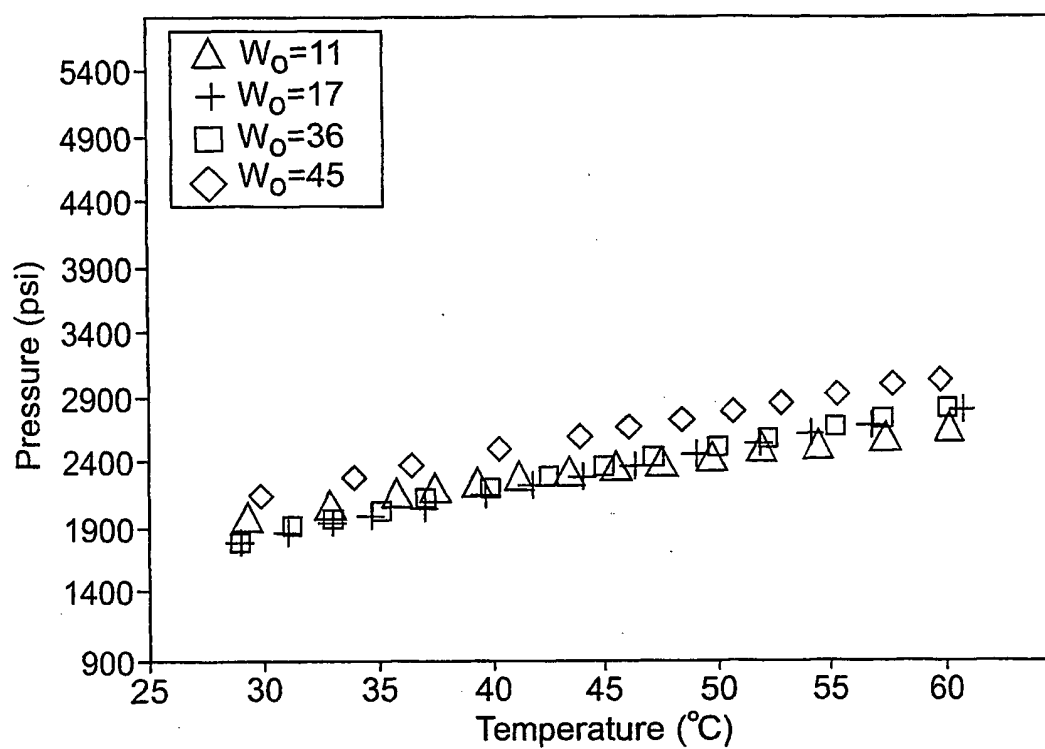
$$W_0 = \frac{\text{moles H}_2\text{O}}{\text{moles surf.}}$$
FIG. 7.

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2.5 wt.% in CO₂

$$W_0 = \frac{\text{moles H}_2\text{O}}{\text{moles surf.}}$$
FIG. 8.

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2.5 wt.% in CO_2

$$W_0 = \frac{\text{moles H}_2\text{O}}{\text{moles surf.}}$$
FIG. 9.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US01/29812

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : C11D 1/02, 1/62, 3/24, 3/36, 7/36; DO6L 1/02, 1/04, 1/22
US CL : 510/175, 176, 285, 412; 8/142
According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 510/175, 176, 285, 412; 8/142

Documentation searched other than minimum documentation to the extent that such documents are included in the fields
~~searched~~

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

WEST

search terms: reverse micelle, supercritical carbon dioxide, surfactant, fluorocarbon, phosphate

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5,676,705 A (JURELLER et al) 14 October 1997, See Abstract; See col. 5, line 1 to column 20, line 60.	1-69
X	US 5,683,473 A (JURELLER et al) 04 November 1997, See Abstract; column 3, line 60 to column 18, line 65.	1-69
X	US 6,114,295 A (MURPHY) 05 September 2000, See Abstract; col. 4, line 5 to col. 8, line 45.	1-69
X	US 5,858,022 A (ROMACK et al) 12 January 1999, See Abstract; column 2, line 30 to column 4, line 60.	1-69

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
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"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"G" document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search
14 DECEMBER 2001

Date of mailing of the international search report
03 JAN 2002

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